AMENDMENTS TO THE CLAIMS

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- 1. (Withdrawn) A method of screening a helminthic parasite preparation that alters a regulatory T cell activity, said method comprising the steps of:
 - (a) obtaining a helminthic parasite preparation;
 - (b) contacting said helminthic parasite preparation with a target; and
- (c) determining the level of an internal marker for regulatory T cell activity in said target after said contacting, wherein a change in said level of said internal marker after said contacting is indicative of said helminthic parasite preparation altering a regulatory T cell activity.
- 2. (Withdrawn) The method of claim 1, wherein said internal marker is a transcription factor.
- 3. (Withdrawn) The method of claim 2, wherein said transcription factor is Scurfin, Smad7, Gata3, or Tbet (Tbx21).
- 4. (Withdrawn) The method of claim 2, wherein said level of said transcription factor is measured at its protein or mRNA level.
- 5. (Withdrawn) A method of screening a helminthic parasite preparation that alters a regulatory T cell activity, said method comprising the steps of:
 - (a) obtaining a helminthic parasite preparation;
 - (b) contacting said helminthic parasite preparation with a target; and
- (c) determining the level of a cell surface marker for regulatory T cell in said target after said contacting, wherein a change in said level of said cell surface marker after said contacting is indicative of said helminthic parasite preparation altering a regulatory T cell activity.

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- 6. (Withdrawn) The method of claim 5, wherein said cell surface marker is selected from the group consisting of: CD4, CD45RB lo , CD45Rc, Cytplytic T lymphocyte associated antigen 4 (CTLA-4), Ox40, 4-1BB, CD25, CD103, CD62L, $\alpha_{E}\beta$ integrin, latency-associated peptide (LAP) or glucocorticoid induced TNF receptor family related protein (GITR), chemokine receptor CCR5, TI-ST2.
- 7. (Withdrawn) The method of claim 6, wherein said level of said surface marker is measured at it protein or mRNA level.
- 8. (Currently amended) A method for treating an animal with a Th1 or Th2 related disease by administering a helminthic parasite preparation that alters a regulatory T cell activity to said animal; and determining the level of regulatory T cell activity, wherein an increase in regulatory T cell activity after said administering is indicative of successful treatment.
- 9. (Withdrawn) A method for monitoring the treatment efficacy of a helminthic parasite preparation for an autoimmune or allergy disease in an animal comprising:
- (a) administering a composition comprising a helminthic parasite preparation or a fraction thereof to said animal; and
- (b) determining the level of a regulatory T cell activity in said animal after said administering, wherein an increase in said level of said regulatory T cell activity after said administering is indicative of the treatment efficacy of said helminthic parasite preparation.

10-16. (Cancelled)

- 17. (New) The method of claim 8, wherein said regulatory T cell activity is measured by determining the level of a regulatory T cell marker.
- 18. (New) The method of claim 17, wherein said regulatory T cell marker is an internal marker.

19. (New) The method of claim 18, wherein said internal marker is Scurfin, Smad7, Gata3, or Tbet (Tbx21).

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- 20. (New) The method of claim 17, wherein said regulatory T marker is a cell surface marker.
- 21. (New) The method of claim 20, wherein said cell surface marker is selected from the group consisting of: CD4, CD45RB^{lo}, CD45Rc, Cytotoxic T lymphocyte associated antigen 4 (CTLA-4), Ox40, 4-1BB, CD25, CD103, CD62L, $\alpha_E\beta$ integrin, latency-associated peptide (LAP) or glucocorticoid induced TNF receptor family related protein (GITR), , chemokine receptor CCR5, TI-ST2.
- 22. (New) The method of claim 17, wherein said regulatory T cell marker is a secreted marker.
- 23. (New) The method of claim 22, wherein said secreted marker is IL4, IL13, IL-5, IL-10 or TGFβ, PgE2.